

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11) EP 0 948 956 A2

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication: 13.10.1999 Bulletin 1999/41

(51) Int Cl.6: A61K 6/083

- (21) Application number: 99106331.4
- (22) Date of filing: 26.03.1999
- (84) Designated Contracting States:

 AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU

 MC NL PT SE

 Designated Extension States:

 AL LT LV MK RO SI
- (30) Priority: 27.03.1998 US 79777 P
- (71) Applicant: DENTSPLY DETREY GmbH D-63303 Dreieich (DE)
- (72) Inventors:
 - Klee, Joachim E., Dr. 78315 Radolfzell (DE)

- Walz, Uwe 78467 Konstanz (DE)
- Lu, Kawang
 Dover, DE 19901 (US)
- (74) Representative: Hartz, Nikolai F., Dr. et al Wächtershäuser & Hartz, Patentanwälte, Tal 29 80331 München (DE)

- (54) Self-curing dental adhesive
- (57) A self-curing dental adhesive has at least one polymerizable resin; at least one polymerizable monomer; at least one polymerizable monomer with an acidic moiety selected from the group

and salts thereof; a stabilizer; an organic solvent or water in a content of from about 10 to about 90 percent by weight; and, at least one part of a thermal redox initiator system or a photoinitiator. The adhesive polymerizes immediately after application of a self-curing dental material that comprises a redox-initiator system, and has an adhesion to dentine of at least about 7 MPa.

BNSDOCID: <EP

Description

5

10

15

20

25

30

35

40

45

50

Technical Field

[0001] The invention concerns a self-curing dental adhesive, which polymerizes after application of a self-curing dental restorative material that comprises a metal-based redox-initiator system without mixing of both systems.

Background of the Invention

[0002] Presently, the dental treatment of cavities of the teeth usually includes using dental composites, compomers or glass ionomers. The use of dental composites and compomers is connected with the application of the adhesive technique. Commonly, both adhesive and dental restorative materials are cured by irradiation with visible light. Dental composites have numerous advantages: high mechanical strength, excellent esthetic properties and a high abrasion resistance. These positive properties are complemented by the fluoride releasing properties of compomers.

[0003] The use of glass ionomers is advantageous at least in part because they are self-adhesive, biocompatible and usable without requiring irradiation by light. Furthermore, they can release fluoride ions. Otherwise, glass ionomers have a relatively low flexural strength and a high abrasion.

[0004] In order to develop a self-curing restorative material with improved mechanical properties, high abrasion resistance and fluoride release a thermal setting adhesive is required.

[0005] Recently, a dual-curing adhesive comprising sulfinates and champhor quinone/amines as polymerization initiator was developed by Dentsply Caulk, Milford DE.

[0006] Generally, a self-curing material is composed of at least of two components, due to the fact that the parts of the initiator system must be distributed in these components.

Summary of the Invention

[0007] It is an object of the present invention to provide a dental adhesive material.

[0008] It is another object of the invention to provide such a material that is self-curing.

[0009] These and other objects of the invention which shall become apparent from the present discussion and the following specification are accomplished by the invention as hereinafter described and claimed.

[0010] In general, a self-curing dental adhesive, comprises:

- (i) at least a polymerizable resin
- (ii) at least a polymerizable monomer
- (iii) at least a polymerizable monomer with an acidic moiety selected from the group consisting of:

and salts thereof

- (iv) at least a stabilizer,
- (v) at least an organic solvent or water in a content of 10 to 90 percent weight of the adhesive and,
- (vi) at least one part of a thermal redox initiator system or a photoinitiator,

wherein the dental adhesive is polymerizable immediately after application of a self-curing dental material that comprises a redox-initiator system, and wherein the dental adhesive has an adhesion to dentine of at least about 7 MPa.

Preferred Embodiments for Carrying Out the Invention

[0011] The present invention includes a self-curing dental adhesive that comprises at least a polymerizable resin, a polymerizable monomer, a polymerizable monomer with at least an acidic moiety selected from the group

55

or salts thereof, a stabilizer, an organic solvent and/or water in a content of 10 to 90 percent, and at least one part of a thermal redox initiator system and/or a photoinitiator that polymerizes immediately after application of a self-curing dental material that comprises a redox-initiator system.

[0012] The adhesion of the invented self-curing adhesive is at least about 7 MPa, preferably 9 MPa, most preferably 11 MPa.

[0013] Preferably, the self-curing dental adhesive contains a salt of a metal or an organo-metalic compound, an amine or a peroxide. Usable metal salts or organo-metalic compounds are selected from the group consisting of copper, silver, cerium, iron, chromium, nickel, vanadium and manganese. Frequently, the metal containing compound is an acetate, salicylate, naphenoate, thiourea complex, acetylacetonate or ethylene tetramine acidic acid.

[0014] The preferred amines are alkyl aryl amines, dialkyl aryl amines, trialkyl amines or derivatives therefrom.

[0015] Preferred peroxides include for example, diacyl peroxides, peresters, perketales, peroxy dicarbonates, dialkyl peroxides, ketone peroxides or alkyl hydroxyperoxides, such as benzoylperoxide, 2,5-dimethyl-2,5-di(benzolyperoxy) hexane, tert.-butylamyl peroxide, di-(tert.-butyl) peroxide, cumenhydoperoxide, tert.-butylhydroperoxide, tert.-butylperoxy-(3,5,5-tri-methylhexanoate) and tert.-butylperoxy-2-ethylhexyl carbonate, laurylperoxide.

[0016] For example a dental adhesive comprising cupric acetylacetonate is applied in a cavity. The cavity is then filled by using a thermal setting material, that comprises cupric thiourea. Both materials polymerize together, whereby the self-curing restorative material also initiates the polymerization of the bonding material. The adhesion of this adhesive/restorative material is 9.0 ± 1.5 MPa.

Comparative Example 1

Dental adhesive

5

10

15

20

25

30

35

45

[0017] Equal amounts of Prime & Bond 2.1™ (Dentsply, De Trey, Constance) and the Self Cure Activator (Dentsply, Caulk, Milford DE) were mixed.

Self-curing dental restorative material

[0018] The used dental restorative material is composed of a powder and a liquid. These have the following composition:

Powder:

[0019] 50.3639 g silylated Strontium-alumo-silicate glass, 2.1546 g silylated Strontium-alumo-silicate glass comprising 2 wt-% cupric thiourea complex and 1.0773 g silylated Strontium-alumo-silicate glass containing 10 wt-% ascorbic acid were mixed homogeneously.

Liquid:

[0020] 18.0000 g of dipentaerthrytrolpentaacrylate monophosphate and 2-(dimethyl) aminoethyl methacrylate (AP-1) comprising 30 wt-% diethylene glycol dimethacrylate. 4.5000 g of macromonomer M-1 (synthesized by reaction of two mols of methacrylic acid, two mols of 2,2-bis-[4-(2,3-epoxypropoxy)-phenyl]-propane and one mole of adipic acid) comprising 30 wt-% diethylene glycol dimethacrylate), 10.6328 g of diethylene glycol dimethacrylate, 0.0167 g of N, N-bis(β-hydroxyethyl)-p-toluidine, 0.5049 g of tert.-butyl peroxy benzoate, 0.0113 g of 2,6-di-tert.-butyl-p-cresol were mixed homogeneously.

Application

[0021] Immediately before use the dental adhesive is applied. Thereafter the restorative material obtained by homogeneous mixture of powder and liquid, in the weight (wt.)-ratio 3.2:1.0, was applied. The working time is 3.75 minutes and the setting time is 3.00 minutes. The adhesion to dentine was measured to be 8.58 ± 1.32 MPa. The dental restorative material shows the following mechanical properties: compressive strength of 256.6 ± 11.9 MPa, a flexural

strength of 87.0 \pm 3.4 and an E-modules of 7780 \pm 785 MPa.

Example 1

10

20

30

35

45

5 Dental adhesive A

[0022] 3.2419 g of 7,7,9,63,63,65-hexamethyl-4,13,60,69-tetraoxo-3,14,19,24,29,34,39,44,49,54, 59,70-dodecane-oxa-5,12,61,68-letraaza doheptaconta diyl -1,72- dimethacrylate (R 5621), 3.7406 g of dipentaerythrol pentaacrylate monophosphate, 8.2819 g of 2,7,7,9,15-pentamethyl-4,13-dioxo-3,14-diox-5,12-diaza-hexadecanediyl-1,16-dimethacylate, 0.4417 g of bisphenol-A-dimethacrylat, 0.1496 g of cetylaminhydrofluorid, 0.0377 g of 2.6-di-tert.-butyl-p-cresol and 0.0330 g of cupric-II-acetylacetonate were dissolved in 59.9736 g of acetone.

Self-curing dental restorative material

[0023] The dental restorative material is composed of powder and liquid as described in comparative example 1.

Application

[0024] Immediately before use the dental adhesive is applied. Thereafter the restorative material obtained by homogeneous mixed of powder and liquid, in the wt.-ratio 3.2:1.0, was applied. The working time is 3.75 minutes and the setting time is 3.00 minutes. The adhesion to dentine was measured to be 9.0 ± 1.5 MPa. The dental restorative material shows the following mechanical properties: compressive strength of 256.6 \pm 11.9 MPa, a flexural strength of 87.0 \pm 3 4 and an E-modules of 7780 \pm 785 MPa.

25 Example 2

Dental adhesive A

[0025] 4.7593 g dipentaerythrol pentaacrylate monophosphate. 7.7083 g 2,7,7,9,15-pentamethyl-4,13-dioxo-314-diox-5,12-diazahexadecanediyl-1,16-dimethacylate, 2.2130 g of 7,7,9,63,63,65-hexamethyl-413.60,69-tetraoxo-3,14,19,24,29,34,39,44,49,54,59,70-dodecaneoxa-5,12,61,68-tetraaza doheptacontadiyl-172-dimethacrylate (R 5621), 1.3177 g trimethylolpropane triacrylate, 2.8213 g diethylene glycol dimethacrylate, 0.0997 g cuppric acetyl acetonate, 0.1922 g 2,6-di-tert.-butyl-p-cresol, 0.1922 g cetylaminhydrofluorid and 1.6952 g of a nanofiller (av. particle size 7 nm), were dissolved in 79.0011 g of acetone.

Self-curing dental restorative material

[0026] The dental restorative material is composed of powder and liquid as described in comparative example 1.

40 Application

[0027] Immediately before use the dental adhesive is applied. Thereafter the restorative material obtained by homogeneous mixed of powder and liquid, in the wt.-ratio 3.4:1.0, was applied. The working time is 3.25 minutes and the setting time is 3.00 minutes. The adhesion to dentine was measured to be $\underline{11.1 \pm 1.8 \text{ MPa}}$.

Example 3

Dental adhesive A

[0028] 4.7137 g dipentaerythrol pentaacrylate monophosphate, 7.6345 g 2,7,7,9,15-pentamethyl-4,13-dioxo-3,14-diox-5,12-diazahexadecanediyl-1,16-dimethacylate, 2.1918 g of 7,7,9,63,63,65-hexamethyl-4,13,60,69-tetraoxo-3,14,19,24,29,34,39,44,49,54, 59,70-dodecaneoxa-5,12,61,68-tetraaza doheptacontadiyl-1,72-dimethacrylate (R 5621), 1.3051 g trimethylolpropane triacrylate, 2.7943 g diethylene glycol dimethacrylate, 1.0559 g p-dimethyl benzoic acid ethylester, 0.1904 g 2,6-di-tert.-butyl-p-cresol, 0.1904 g cetylaminhydrofluorid and 1.6790 g of a nanofiller (av. particle size 7 nm), were dissolved in 78.2450 g of acetone.

Self-curing dental restorative material

[0029] The dental restorative material is composed of powder and liquid as described in comparative example 1.

5 Application

10

20

25

30

35

40

45

50

55

[0030] Immediately before use the dental adhesive is applied. Thereafter the restorative material obtained by homogeneous mixed of powder and liquid, in the wt.-ratio 3.4:1.0, was applied. The working time is 3.25 minutes and the setting time is 3.00 minutes. The adhesion to dentine was measured to be 8.9 ± 2.6 MPa.

Example 4

Dental adhesive A

[0031] 3.2419 g of 7,7,9,63,63,65-hexamethyl-4,13,60,69-tetraoxo-3,14,19,24,29,34,39,44,49,54, 59,70-dodecane-oxa-5,12,61,68-tetraaza doheptacontadiyl -1,72- dimethacrylate (R 5621), 3.7406 g of dipentaerythrol pentaacrylate monophosphate, 8.2819 g of 2,7,7,9,15-pentamethyl-4,13-dioxo-3,14-diox-5,12-diaza-hexadecanediyl-1,16-dimethacrylate, 0.4417 g of bisphenol-A-dimethacrylat, 0.1496 g of cetylaminhydrofluorid, 0.0377 g of 2,6-di-tert.-butyl-p-cresol and 0.2283 g of cupric-II-acetylacetonate were dissolved in 59.9736 g of acetone.

Self-curing dental restorative material

.Powder:

[0032] 89.10 g silylated Strontium-alumo-silicate glass, 10-10 g OX-50 organic filler, 1.00 g Aerosil R-972, 0.02 g cupric acetylacetonate and 0.15 g ascorbyl palmitate were mixed homogeneously.

Liquid:

[0033] 10.00 g OEMA, 10.00 g dipentaerthrytrolpentaacrylate monophosphate, 30.00 g diethylene glycol dimethacrylate, 50.00 g 2,2-bis-[p-(2-hydroxy-3-methacryloyloxypropoxy)-pheny]-propane, 2.00 g distilled water and 0.20 g tert.-butyl peroxy benzoate were mixed homogeneously.

[0034] Apply one coat of the primer to etched dentine, air dry. Triturate powder and liquid of the restorative in capsule at 3.75.1 ratio for 12 seconds. Apply the paste to dentine in a post. Let the assembly self cure at 37°C. Then stored in 37°C water for 24 hours. Test sheer bond strength on Instron. The average bond strength was 9.18 Mpa.

[0035] It is evident therefore, that the objects of a a self-curing dental adhesive are carried out by the invention as herein described. All possible aspects of the invention beyond the best mode have not been necessarily described, and the scope of the invention shall only be determined by the following claims.

Claims

- 1. A self-curing dental adhesive, comprising
 - (i) at least one polymerizable resin;
 - (ii) at least one polymerizable monomer;
 - (iii) at least one polymerizable monomer with an acidic moiety selected from the group consisting of

and salts thereof;

- (iv) a stabilizer;
- (v) an organic solvent or water in a content of from about 10 to about 90 percent; and,
- (vi) at least one part of a thermal redox initiator system or a photoinitiator;

wherein the adhesive polymerizes immediately after application of a self-curing dental material that comprises a redox-initiator system, and has an adhesion to dentine of at least about 7 MPa.

- 2. A self-curing dental adhesive of claim 1 having an adhesion to dentine of at about 11 MPa.
- 3. A self-curing dental adhesive of claim 1 wherein said thermal redox initiator system is a salt of a metal or an organometalic cutaining compound, an amine or a peroxide.
- A self-curing dental adhesive of claim 3 wherein said metal of said metal containing compound is selected from the group consisting of copper, silver, cerium, iron, chromium, nickel, vanadium and manganese.
 - A self-curing dental adhesive of claim 3 wherein said metal containing compound is an acetate, salicylate, naphenoate, thiourea complex, acetylacetonate or ethylene tetramine acidic acid.
- 6. A self-curing dental adhesive of claims 3 wherein said amine is is an alkyl aryl amine, a dialkyl aryl amine, a trialkyl amine or derivatives therefrom.
 - 7. A self-curing dental adhesive of claims 3 wherein said peroxide is a diacyl peroxide, a perester, a perketale, a peroxy dicarbonate, a dialkyl peroxide, a ketone peroxide or a alkyl hydroxyperoxide.
 - 8. A self-curing dental adhesive of claims 3 wherein said peroxide selected from benzoylperoxide, 2,5-dimethyl-2,5-di (benzolyperoxy)hexane, tert.-butylamyl peroxide, di-(tert.-butyl) peroxide, cumenhydoperoxide, tert.-butylhydroperoxide, tert.-butylperoxy-(3,5,5-tri-methylhexanoate) and tert.-butylperoxy-2-ethylhexyl carbonate, laurylperoxide.

6

5

20

25

30

35

40

45

50

55

Europäisches Patentamt European Patent Office Office européen des brevets



(11) EP 0 948 956 A3

(12)

EUROPEAN PATENT APPLICATION

(88) Date of publication A3: 19.07.2000 Bulletin 2000/29

(51) Int Cl.7: A61K 6/083

- (43) Date of publication A2: 13.10.1999 Bulletin 1999/41
- (21) Application number: 99106331.4
- (22) Date of filing: 26.03.1999
- (84) Designated Contracting States:

 AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU

 MC NL PT SE

 Designated Extension States:

 AL LT LV MK RO SI
- (30) Priority: 27.03.1998 US 79777 P
- (71) Applicant: DENTSPLY DETREY GmbH D-63303 Dreieich (DE)
- (72) Inventors:
 - Klee, Joachim E., Dr. 78315 Radolfzell (DE)

- Walz, Uwe 78467 Konstanz (DE)
- Lu, Kawang Dover, DE 19901 (US)
- (74) Representative: Hartz, Nikolai F., Dr. et al Wächtershäuser & Hartz, Patentanwälte, Tal 29 80331 München (DE)

- (54) Self-curing dental adhesive
- (57) A self-curing dental adhesive has at least one polymerizable resin; at least one polymerizable monomer, at least one polymerizable monomer with an acidic moiety selected from the group

and salts thereof; a stabilizer; an organic solvent or water in a content of from about 10 to about 90 percent by weight; and, at least one part of a thermal redox initiator system or a photoinitiator. The adhesive polymerizes immediately after application of a self-curing dental material that comprises a redox-initiator system, and has an adhesion to dentine of at least about 7 MPa.



EUROPEAN SEARCH REPORT

Application Number EP 99 10 6331

·	DOCUMENTS CONSIDERE Citation of document with indica	tion, where appropriate,	Relevant	CLASSIFICATION OF THE APPLICATION (Int.Cl.5)	
ategory	of relevant passages		to claim		
(EP 0 218 248 A (MUEHLB 15 April 1987 (1987-04 * column 4, line 19 - * column 15, line 50 -	line 56 * - column 16, line 22	1,3-8	A61K6/083	
	* column 17, line 1 - * column 17, line 50 - * claims; examples *	- (ine 50 *			
Α .	US 5 556 897 A (HONDA 17 September 1996 (19 * column 2, line 32 - * column 3, line 10 - * column 3, line 53 - * column 9, line 63 - * column 11, line 11	96-09-17) line 67 * line 35 * column 9, line 57 * column 10, line 13	1-8 *		
A	EP 0 811 368 A (TOKU) 10 December 1997 (1991) * page 5, line 4 - 1 * page 5, line 35 - 1	97-12-10) ine 20 *	1-8	TECHNICAL FIELDS SEARCHED (Int.Cl.5)	
А	WO 96 19179 A (DENTS 27 June 1996 (1996-0 * page 13, paragraph paragraph 1 * * application exampl	6-27) 13 - page 17,	1-8	A61K	
A	EP 0 712 621 A (HERA 22 May 1996 (1996-05 * page 3, line 8 - 1	5-22)			
A	GB 2 261 223 A (G C 12 May 1993 (1993-0! * page 3, paragraph 3 *	DENTAL IND CORP) 5-12) 2 - page 9, paragrap	1-8		
		to all daims			
The present search report has Proce of search THE HAGUE CATEGORY OF CITED DOCUMENT		Date of completion of the search		Examiner	
		23 May 2000		Cousins-Van Steen, G	
AM 1503 03 82	CATEGORY OF CITED DOCUMENTS X : particularly resevant if taken alone Y : particularly resevant if combined with and cocument of the same category A : technological background O : non-written disclosure P : intermediate document	E : earlier par after the fi other D : document L : document	ing date in the app in cited in the app in cited for other report the same pate	ication	



EUROPEAN SEARCH REPORT

EP 99 10 6331

	DOCUMENTS CONSIDERE				
Category	Citation of document with indicate of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.CI.8)		
Ρ,Χ	WO 98 46198 A (MINNESOT 22 October 1998 (1998-1 * page 1, line 16 - lin * page 12, line 8 - lin * page 12, line 17 - li * page 13, line 21 - pa * page 17, line 3 *	0-22) ne 28 * ne 11 * ne 23 *	1,3-8		
:					
				TECHNICAL FIELDS SEARCHED (Int.Cl.6)	
	The present search report has been o	drawn up for all claims			
	Place of search	Date of completion of the search	Date of completion of the search Examiner		
	THE HAGUE	23 May 2000	Cou	Cousins-Van Steen, G	
X : par Y : par doo A : teo	CATEGORY OF CITED DOCUMENTS ritoularly relevant if taken alone riticularly relevan; if combined with another ament of the same category hnological background n-written disclosure	T : theory or princi E : earlier patent d after the timing o D : document ofter L : document circle	ple underlying the locument, but publi tate if in the application if for other reasons	invention ished on, or	

3

ANNEX TO THE EUROPEAN SEARCH REPORT ON EUROPEAN PATENT APPLICATION NO.

EP 99 10 6331

1 1 m

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

23-05-2000

Patent document cited in search report		Publication date			Publication date	
	Α	15-04-1987	DE	3536077 A	09-04-1987	
EP 0218248	A	19-04-1907	JP	2588702 B	12 -03-199 7	
				62149707 A	03-07-1987	
			üs	4806381 A	21-02-1989	
 US 5556897	Α	17-09-1996	JP	6024928 A	01-02-1994	
EP 0811368	Α	10-12-1997	US	5866631 A	02-02-1999	
FL 0911300	^	10 12 1337	WO	9723191 A	03-07-1997	
			JP	9227325 A	02-09-1997	
	Α	 27-06-1996	AU	695605 B	20-08-1998	
WO 9619179	A	27-00 1930	ΑŬ	4425696 A	10-07-1996	
			BR	9510062 A	02-06-1998	
			CA	2202755 A	27-06-1996	
			EP	0799016 A	08-10-1997	
			JΡ	10511354 T	04-11-1998	
			ZA	9510792 A	20-06-1996	
50.0712621		22-05-1996	DE	4441124 A	23-05-1996	
EP 0712621	Α.	22 03 1330	CA	2163166 A	19-05-1996	
			CN	1129150 A	21-08-1996	
			JP	8224535 A	03-09-1996	
			US	5618372 A	08-04-1997	
		12-05-1993	DE	4237391 A	13-05-199	
GB 2261223	A	17-02-1337	JP	5186309 A	27-07-1 99 3	
			ÜS	5290172 A	01-03-199	
10.0046169		22-10-1998	US	5922786 A	13-07-199	
WO 9846198	A	25-10-1330	AU	5179998 A	11-11-199	
			EP	0973482 A	26-01-200	

© For more details about this annex :see Official Journal of the European Patent Office, No. 12/82